The solid pharmaceutical formulations of this invention may comprise any conventional solid dosage form including, for example, tablets, pills, capsules, and the like. The formulations may further comprise a binder such as hydroxypropylmethyl cellulose, gum tragacanth, acacia gum, corn starch or gelatin; a disintegrating agent such as corn starch, potato starch, alginic acid, sodium starch gylcolate, croscamellose sodium, or crospovidone; and a sweetening agent such as sucrose, lactose, or saccharin. If desired, the formulations may further comprise_lyoprotectants, including, for example, sugars and amino acids, buffers, and stabilizers. Various other materials may be present in the form of coatings or to modify the physical form of the dosage unit. For instance, tablets may be coated with mixtures comprising, for example, titanium dioxide, dextrose, polyethylene glycol, sodium carboxymethyl cellulose, dextrin, and the like. The coatings may also comprise the form of an enteric polymer including phthalate derivatives, such as cellulose acetate phthalate, polyvinylacetate phthalate and hydroxypropylmethyl cellulose phthalate, polyacrylic acid derivatives, such as methacrylic acid copolymer, vinyl acetate, and crotonic acid copolymers.